LISTING OF CLAIMS:



- 1. (Canceled)
- 2. (Canceled)
- 3. (Canceled)

4. (ORiginal). A piperazinylalkylbenzofuran derivative of the formula

as claimed in Claim 1, wherein represents a C₁₋₄ alkyl group, ${ t R}^2$ stands for a hydrogen atom, means an oxygen atom, is a hydroxy group, Z represents a hydrogen atom, Ar' represents a diphenylmethyl group, a pyridyl group, a partially saturated 5-membered heterocyclic group containing two oxygen_atoms and being condensed with a phenyl group, or a phenyl group substituted by substituents Ro, Ro and R^7 , wherein R^5 , R^6 and R^7 mean, independently, a hydrogen atom, a halo atom, a trifluoromethyl group, a C1_4 alkyl group, a C_{1-4} alkoxy group, or a methylenedicxy group,

n has a value of O or 1, and pharmaceutically suitable acid addition salts thereof.

5. (Original) A piperazinylalkylbenzofuran derivative as claimed in Claim 4, wherein in formula Ia R¹ represents a methyl group, R² stands for a hydrogen atom, X means an oxygen atom, is a hydroxy group, z represents a hydrogen atom, Ar' represents a diphenylmethyl group, a pyridyl group, a benzo-1,3-dioxolanyl group or a phenyl group optionally substituted by one or two halo atom(s), one or two methyl group(s), a methylenedioxy group, a trifluoromethyl group or a methoxy group, n has a value of O or 1, and pharmaceutically suitable acid addition salts thereof.

- 7. (Original). A process for the preparation of a benzofuran derivative of the formula I, wherein R¹, R², Z, X, Y, A, B and Ar are as defined in Claim 1, or a pharmaceutically suitable acid addition salt thereof, characterized in that
 - a) a halide of the formula

wherein \mathbb{R}^1 , \mathbb{R}^2 , X, Y and Z are as defined in connection with formula I, Hal represents a halo atom, is reacted with a secondary amine of the formula

wherein A, B and Ar are as stated in connection with formula I; or

b) for the preparation of a benzofuran derivative of the formula I, wherein Y represents a hydroxy group, R^1 , R^2 , X, Z, A, B and Ar are as defined in connection with formula I, an epoxide of the formula

wherein R^1 , R^2 , Z and X are as defined above, is reacted with a secondary amine of the

wherein R^2 , A, B and Ar are as stated above; or

- e) for the preparation of a benzofuran derivative of the formula I, wherein A forms with B a group of the formula -C=C-, R¹, R², X, Y, Z and Ar are as defined in connection with formula I, a benzofuran derivative of the formula I, wherein A stands for a group of the formula COH, B represents a methylene group, R¹, R², X, Y, Z and Ar are as stated above, is dehydrated; or
- f) for the preparation of a benzofuran derivative of the formula I, wherein A represents a group of the formula COH, B stands for a methylene group, R¹, R², X, Y, Z and Ar are as defined in connection with formula I, however, Ar is other than a hydrogen atom, a ketone of the formula

wherein R^1 , R^2 , X, Y and Z are as stated above, is reacted with an arylmagnesium halide of the formula

Hal-Mg-Ar

XVI

wherein Ar is as stated above, Hal represents

a halo atom, and the adduct formed is decomposed with water; or

g) for the preparation of a benzofuran derivative of the formula I, wherein A represents a group of the formula COH, B stands for a methylene group, R¹, R², X, Y, Z and Ar are as defined in connection with formula I, but Ar is other than a hydrogen atom, a ketone of the formula XV, wherein R¹, R², X, Y and Z are as stated above, is reacted with an aryl lithium compound of the formula

Li-Ar

IIVX

wherein Ar is as stated above, and the adduct formed is decomposed with water; or

- h) for the preparation of a benzofuran derivative of the formula I, wherein A represents a group of the formula CH, B stands for a methylene group, R¹, R², X, Y, Z and Ar are as defined in connection with formula I, a compound of the formula I, wherein A forms with B a group of the formula -C=C-, R¹, R², X, Y, Z and Ar are as stated above, is hydrogenized; or
- i) for the preparation of a benzofuran derivative of the formula I, wherein A represents a group of the formula CH, B stands for a methylene group, R^1 , R^2 , X, Y, Z and Ar are as defined in connection with formula I, an epoxide of the formula III, wherein R^1 , R^2 , Z and X are as stated above, is reacted

with a secondary amine of the formula IV, wherein A stands for a group of the formula CHOH, B and Ar are as stated above, under dehydrating reaction conditions, and the formed compound of the formula I, wherein A forms with B a group of the formula -C=C-, R¹, R², X, Y, Z and Ar are as stated above, is hydrogenized in the reaction mixture in which it was prepared; and

if desired, an obtained base of the formula I is reacted with an inorganic or organic acid to form a pharmaceutically suitable acid addition salt thereof, or liberated from the acid addition salt with a base.

- 8. (Canceled)
- 9. (Canceled)
- 10. (Canceled)

11. (Original). A pharmaceutical composition as claimed in Claim 8, comprising a piperazinylalkylbenzofuran derivative of the formula

$$R^{\frac{1}{N}} = R^{\frac{N}{N}} =$$

wherein

 R^{1} represents a C_{1-4} alkyl group, R^{2} stands for a hydrogen atom,

X means an oxygen atom,

Y is a hydroxy group,

Z represents a hydrogen atom,

Ar' represents a diphenylmethyl group, a pyridyl group, a partially saturated 5-membered heterocyclic group containing two oxygen atoms and being condensed with a phenyl group, or a phenyl group substituted by substituents R⁵, R⁶ and R⁷, wherein

 R^5 , R^6 and R^7 mean, independently, a hydrogen atom, a halo atom, a trifluoromethyl group, a C_{1-4} alkyl group, a C_{1-4} alkoxy group, or a methylenedioxy group,

n has a value of O or 1, or a pharmaceutically suitable acid addition salt thereof as the active ingredient.

12. (Original). A pharmaceutical composition as claimed in Claim 11, comprising a piperazinylalkylbenzofuran derivative of the formula Ia, wherein

R represents a methyl group,

R² stands for a hydrogen atom,

x means an oxygen atom,

Y is a hydroxy group,

z represents a hydrogen atom,

Ar' represents a diphenylmethyl group, a pyridyl group, a benzo-1,3-dioxolanyl group or a phenyl group optionally substituted

by one or two halo atom(s), one or two
methyl group(s), a methylenedioxy group,
a trifluoromethyl group or a methoxy group,
n has a value of 0 or 1,
or a pharmaceutically suitable acid addition
salt thereof as the active ingredient.

- 13. (Canceled)
- 14. (Canceled)
- 15. (Canceled)

16. (Original) A halide of the formula

$$R \downarrow 0$$
 $X \downarrow Hal$ R^2

wherein

 R^1 and R^2 represents, independently, a hydrogen atom or a C_{1-4} alkyl group,

X stands for an oxygen atom or a sulfur atom,

Y means a hydrogen atom or a hydroxy group,

Z represents a hydrogen atom, a halo atom, a C_{1-4} alkyl group, a C_{1-4} alkoxy group, an amino group, a nitro group, a cyano group, a trifluoromethyl group or a group of the formula $-\text{COOR}^3$, $-\text{NHCOR}^3$ or $-\text{SO}_2\text{NR}^3\text{R}^4$, wherein

 ${\ensuremath{\mathbb{R}}}^3$ stands for a hydrogen atom or a ${\ensuremath{\text{C}}}_{1-4}$ alkyl group,

 R^4 means a C_{1-4} alkyl group, or

R³ and R⁴ form, together with the adjacent nitrogen atom, a saturated or unsaturated heterocyclic group having 5 to 10 members and optionally comprising one or more nitrogen atom(s) and/or one or more oxygen atom(s) and/or one or more sulfur atom(s),

Hal represents a halo atom. 17. (Original). A ketone of the formula

$$R \rightarrow 0$$
 $X \rightarrow X$
 $X \rightarrow X$
 $X \rightarrow X$
 $X \rightarrow X$

wherein

- ${\bf R}^1$ and ${\bf R}^2$ represents, independently, a hydrogen atom or a ${\bf C}_{1-4}$ alkyl group,
- X stands for an oxygen atom or a sulfur atom,
- Y means a hydrogen atom or a hydroxy group,
- represents a hydrogen atom, a halo atom, a C_{1-4} alkyl group, a C_{1-4} alkoxy group, an amino group, a nitro group, a cyano group, a trifluoromethyl group or a group of the formula $-COOR^3$, $-NHCOR^3$ or $-SO_2NR^3R^4$. wherein
 - R³ stands for a hydrogen atom or a C₁₋₄ alkyl group,
 - ${\bf R}^4$ means a ${\bf C}_{1-4}$ alkyl group, or ${\bf R}^3$ and ${\bf R}^4$ form, together with the adjacent nitrogen atom, a saturated or unsaturated

heterocyclic group having 5 to 10 members and optionally comprising one or more nitrogen atom(s) and/or one or more oxygen atom(s) and/or one or more sulfur atom(s).